

OXAZOLINE SURFACTANT ANODE ADDITIVE
FOR ALKALINE ELECTROCHEMICAL CELLS

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] Not applicable.

STATEMENT REGARDING FEDERALLY
SPONSORED RESEARCH OR DEVELOPMENT

[0002] Not applicable.

BACKGROUND OF THE INVENTION

[0003] The demand for very small electrochemical cells has increased with increased use of small, electrically-powered devices, and alkaline and alkaline metal-air electrochemical cells are an increasingly popular choice for powering such devices as hearing appliances and computers. Metal air cells contain an anode and an oxygen cathode, with the oxygen usually obtained from ambient air. The oxygen cathode catalytically promotes the reaction of oxygen with an aqueous neutral or alkaline electrolyte and is not consumed or changed during discharge. As the cathode is extremely compact yet has essentially unlimited capacity, very high energy densities are achieved, resulting from the increased available volume for the anode active material. Consequently, a metal-air cell can provide more watt-hours of electromotive force than a so-called "two-electrode cell" of similar cell size, mass and anode composition that contains both anode- and cathode-active materials inside the cell structure. Metal-air cells have an advantageous watt-hour capacity/mass ratio without regard to size or configuration, which can be, e.g., button cells as in, for example, U.S. Patent No. 5,721,065 ("the '065 patent"), or cylindrical cells as in, for example, U.S. Patent No. 6,210,827 (the '827 patent"), each incorporated by reference in its entirety as if set forth herein.

[0004] A typical metal air cell such as a zinc air cell contains an anode that includes a zinc alloy powder, a carbon-based air cathode and an alkaline electrolyte with a gelling agent to prevent the electrolyte from creeping through the cell seals. For efficient operation, the anode of

a zinc air cell should participate only during current-generating reaction periods. In reality, though, corrosive shelf-discharge side reactions in the alkaline electrolyte can reduce both service and shelf life of alkaline electrochemical systems that use zinc as the anode active material. Much effort has been directed to improving electrochemical reaction efficiency and cell output. Previously, mercury was added to cell anodes to improve corrosion resistance. In recent years, mercury has been replaced by substances that conform to environmental requirements. Small amounts of metals such as lead, indium, and bismuth, and combinations thereof, can effectively improve corrosion behavior of anodic zinc. Certain organic surfactants can provide effective corrosion-inhibiting effect at the metal surface while making the anode sufficiently available for electrochemical oxidation that the cell output can be maintained under heavy cell loading. Surfactants can, like mercury, improve discharge capacity, service life and shelf life, without substantial adverse environmental impact. U.S. Patent No. 4,857,424 ("the '424 patent") incorporated by reference herein as if set forth in its entirety, discloses reduced-mercury or mercury-free zinc-manganese dioxide cells containing an organosiliconate type surfactant. Also, the '065 patent discloses a button cell having an anode mix containing zinc metal powder, indium or other compounds, optionally a low level of mercury, a gelling agent and preferably a hydroxyethylcellulose surfactant. The anode disclosed therein sustains longer periods of power production at a relative steady voltage of at least 1.1 volts while protecting the anode metal from corrosion in the alkaline environment.

[0005] Similarly, the '827 patent discloses a cylindrical cell having an anode mix containing electrolyte, a gelling agent, particulate zinc, zinc oxide, additives and an organic surfactant comprising hydroxyethylcellulose.

[0006] Japanese Patent No. JP10083812 to Toshiba Battery Co. Ltd. discloses providing a high performance zinc alkaline battery containing 0.5 – 100 ppm of a fluorine-containing surfactant in a non-amalgamated-zinc-alloy-based gelled anode having a viscosity of 100,000 to 300,000 cPs at 25 °C.

[0007] US Patent No. 5,382,482 concerns suppressing dendrites and shape change in an alkaline cell by disposing a cross-linked polymer film layer in close proximity to the anode active material, where the polymer film layer can contain, among other polymers, polyoxazoline. The patent does not describe including an oxazoline surfactant in the anode mix of the cell.

[0008] Further improvements in performance, such as increased operating voltage, improved discharge profile and reduced sensitivity to open circuit rest in alkaline electrochemical cells, preferably without sacrificing known benefits, are welcome and desired.

BRIEF SUMMARY OF THE INVENTION

[0009] The present invention discloses that an alkaline electrochemical cell having superior discharge performance and corrosion resistance includes an oxazoline surfactant additive in the cell anode. In one aspect, the present invention relates to an anode for use in a alkaline electrochemical cell, where the anode contains a metal anode active material, an anode electrolyte and an oxazoline surfactant. Optionally, the anode of the invention can include additional components known in the manufacture of alkaline electrochemical cells and anodes for same. In a related aspect, the invention relates to an alkaline electrochemical cell containing an anode of the invention. In another aspect, the present invention relates to methods for making an anode of the invention, or for making a cell containing the anode, as are detailed below.

[00010] It is an object of the invention to improve discharge performance and corrosion resistance without sacrificing the known benefits of adding surfactants to alkaline cell anodes.

[00011] It is a feature of the invention that the anode, and an electrochemical cell containing the anode, contain an oxazoline surfactant that can coat at least a portion of the particles of the anode active material.

[00012] It is an advantage of the present invention can maintain known benefits of anode surfactants, and can provide additional benefits that can include improved discharge efficiency at low and high rates, high operating voltage, lack of initial potential dip, and reduced sensitivity to open circuit rest.

[00013] Other objects, features and advantages of the present invention will become apparent in view of the following detailed description of the invention and the drawings.

BRIEF DESCRIPTION OF THE DRAWINGS

[00014] Fig. 1 compares the discharge profiles of cells of the invention containing an oxazoline surfactant and cells containing an organosiliconate surfactant. Discharge was at 374 Ohm for 16 hours/day at 70 °F, 50 % relative humidity, after one month storage at 70°F.

[00015] Fig. 2 compares the discharge profiles of cells of the invention containing an oxazoline surfactant and cells containing an organosiliconate surfactant. Discharge was at 620 Ohm for 16 hours/day at 70 °F 50 % relative humidity, after 20 days storage at 140°F.

DETAILED DESCRIPTION OF THE INVENTION

[00016] The present invention relates to using an oxazoline-type surfactant, which can be a fatty oxazoline surfactant, as an additive for an alkaline electrochemical cell anode. Cells comprising the anode can exhibit improved electrical discharge performance, including higher operating voltage, good high rate pulse capability, no initial potential dip and reduced sensitivity to open circuit rest in comparison to cells that use other surfactants.

[00017] Oxazoline-type surfactants can be added to anode materials of any alkaline metal-anode electrochemical cells to improve discharge performance and to improve discharge capacity, service-life, and shelf life. While the invention was exemplified with metal-air anodes and cells, more particularly in zinc-air cells, the invention can be employed in metal-containing anodes generally, more particularly in zinc anodes, and in cells containing same. Suitable chemistries include, but are not limited to, zinc-air, zinc-manganese dioxide, zinc/AgO and zinc/Ag₂O systems, without regard to size or configuration. Preferable cells are of standard configurations, such as button cells, cylindrical cells and prismatic cells, which are available in a variety of standard sizes. For example, oxazoline type-surfactants can be substituted for the surfactants described in incorporated U.S. Patent Nos. 4,857,424, 5,721,065 and 6,210,827. The anode materials to which an oxazoline surfactant is added contain a metal powder, a gelling agent, and an electrolyte. The anode materials can also include other components including but not limited to mercury, indium, gallium, aluminum, lead, bismuth and calcium compounds, and combinations thereof, as disclosed, for example, in the '065 patent and in US Patent No. 5,721,072, also incorporated herein by reference as if set forth in its entirety.

[00018] In one aspect, the present invention relates to an anode mix that includes a metal powder suitable for an anode of an alkaline electrochemical cell, an electrolyte compatible with the anode metal, an oxazoline surfactant and a gelling agent. In another aspect, the present invention relates to an anode that includes the composition described above in a container for receiving the composition of matter. In still another aspect, the present invention relates to an alkaline electrochemical cell that contains the anode. The cell has a discharge voltage of at least

0.7 volts, preferably 1.0 volts, more preferably 1.1 volts and still more preferably 1.2 volts, during its service life.

[00019] In preferred embodiments, the metal powder is preferably about 55% to about 73% by weight of the anode, the electrolyte is about 27% to about 45% (by weight) in an aqueous solution, and the oxazoline surfactant is about 0.001% to about 5% (by weight), based on the weight of the particles of metal powder. The anode material generally also contains about 0.1% to about 0.5% of gelling agent, by weight based on the weight of active material, and between about 1% to about 4%, preferably about 2% of zinc oxide, by weight based on the weight of the electrolyte. In more preferred embodiments, the metal powder is a particulate zinc alloy powder and the electrolyte is potassium hydroxide.

[00020] In a related aspect, the present invention is a method of making an anode mix for use in the anode and in the cells of the invention, the method involving the steps of mixing a metal powder, a gelling agent, an oxazoline surfactant, and an electrolyte in the above-mentioned proportions. The order in which the components are combined is not critical, except insofar as when mixing the oxazoline surfactant with a liquid (namely, the electrolyte or the gelled electrolyte), the surfactant is preferably a state (e.g., melted). Alternatively, if the surfactant is mixed first with the metal powder or other dry component, it can be a solid. In one embodiment of the method, the gelling agent and the alkaline electrolyte are combined first to form a gelled electrolyte. Then, the oxazoline surfactant is added to the gelled electrolyte with agitation. Finally, the metal powder is combined with the surfactant-containing alkaline electrolyte to form the gelled zinc anode material which can be incorporated into the receiving container.

[00021] In another related aspect, the present invention is a method of making an alkaline electrochemical cell involving the step of incorporating the aforementioned inventive anode into an electrochemical cell, the process for making the cell being, apart from the composition of the anode, conventional as is shown, for example in US Patent No. 4,617,242, incorporated herein by reference as if set forth in its entirety.

[00022] A suitable oxazoline surfactant can be solubilized in an anode-compatible electrolyte and is a liquid or can be liquefied under the anode processing conditions. US Patent No. 3,389,145, incorporated by reference herein as if set forth in its entirety, discloses structures of one suitable set of oxazolines and processes for making same. Also suitable for use in the anode of the invention are substituted oxazoline surfactants having the structures shown in US

Patent No. 3,336,145, in US Patent No. 4,536,300, in US Patent No. 5,758,374 and in US Patent No. 5,407,500, each incorporated by reference herein as if set forth in its entirety, and mixtures of any of the foregoing. A most preferred oxazoline surfactant, ethanol,2,2'-[(2-heptadecyl-4(5H)-oxazolydine) bis (methylenedioxy-2,1-ethanedioxy)] bis, has a structure shown as Formula (I-2) in incorporated US Patent No. 5,407,500. This compound is commercially available from Angus Chemical (Northbrook, IL) as Alkaterge T-IV.

EXAMPLE

[00023] Zinc air cells according to the invention were built with an anode as described in incorporated US Patent No. US Patent No. 4,617,242, except that the oxazoline surfactant was substituted at 0.3 %, relative to the alkaline electrolyte in test cells in place of the organosiliconate surfactant of the prior anode composition. The structure of the test and control cells was as described in incorporated US Patent No. 5,721,065.

[00024] The discharge characteristics of 13 size zinc air cells of the present invention containing Alkaterge T-IV were compared against otherwise identical commercial cells having an organosiliconate surfactant in the anode. Fig. 1 illustrates the discharge curve at 374 Ohm after 1 month storage of cells at 70°F; Fig. 2 shows the discharge curve at 620 Ohm after 20 days storage at 140°F. As shown in Figs. 1 and 2, cells containing oxazoline-type surfactant-based exhibited operating voltage at least 5-10 mV higher than prior art cells during most of their usable lives. In addition, the test cells did not exhibit an initial potential dip and were less sensitive to open circuit rest than otherwise comparable cells containing the organosiliconate type-surfactant.

[00025] Further, Table 1 shows discharge capacity values at 374 Ohm of test cells containing Alkaterge T-IV surfactant. The capacity of these cells was comparable to that of organosiliconate type surfactant-based cells before and after storage for one month at 70°F, 50% relative humidity, although discharge capacity before storage was comparatively lower in cells that contained Alkaterge T-IV.

	Control	Alkaterge T-IV	Alkaterge T	Alkaterge E
No Delay	251	244	247	N/A
One month	252	252	198	N/A

Species	1980	1981	1982	1983	1984	1985	1986	1987	1988	1989	1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017	2018	2019	2020	2021	2022	2023	2024	2025	2026	2027	2028	2029	2030
1980	1981	1982	1983	1984	1985	1986	1987	1988	1989	1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017	2018	2019	2020	2021	2022	2023	2024	2025	2026	2027	2028	2029	2030	

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